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INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Use as many sheets as necessary)				Application Number	10/613,432
				Filing Date	July 3, 2003
				First Named Inventor	Stephen A. Scaringe
				Art Unit	1646
				Examiner Name	Not yet accorded
Sheet	1	of	2	Attorney Docket No.	DH260625 CON2

NON PATENT LITERATURE DOCUMENTS

Examiner Initials*	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s) publisher, city and/or country where published	T ²
Ho	1	WOJCIECH T. MARKIEWICZ, "Tetraisopropylidisiloxane-1, 3-diyl, a Group for Simultaneous Protection of 3' - and 5' - Hydroxy Functions of Nucleosides," J. Chem. Research(s), 1979, pp. 24-25	
Ho	2	SERGE L. BEAUCAGE, et al., "Tetrahedron Report Number 309," Tetrahedron, Vol. 48, No. 12, pp. 2223-2311, 1992	
Ho	3	DANIEL C. CAPALDI, et al., "Use of the 1-(2-fluorophenyl)-4-methoxypiperidin-4-yl (Fpmp) and Related Protecting Groups in Oligoribonucleotide Synthesis: Stability of Internucleotide Linkages to Aqueous Acid," Nucleic Acids Research, 1994, Vol. 22 No. 12, pp. 2209-2216	
Ho	4	CHRIS CHRISTODOULOU, "Incompatibility of Acid-Labile 2' and 5' Protecting Groups For Solid - Phase Synthesis of Oligoribonucleotides," Tetrahedron Letters, 1986, Vol. 27, No. 13, pp. 1521-1522	
Ho	5	TSUJIAKI HATA, et al. "2-Chloroethyl Orthoformate as a Reagent for Protection in Nucleotides Synthesis," Tetrahedron Letters, No. 51, pp. 4443-4446, 1969	
Ho	6	J. GORDON HILL, et al. "Anhydrous tert - Butyl Hydroperoxide in Toluene: The Preferred Reagent for Applications Requiring Dry TBHP," J. Org. Chem., 1983, 48, pp. 3607-3608	
Ho	7	OSAMU ODAI, et al. "Synthesis and NMR Study of Ribooligonucleotides Forming a Hammerhead-type RNA Enzyme System," Nucleic Acids Research, Vol. 18, No. 20, pp. 5955-5960,	1990,
Ho	8	RAO, et al., "Use of the 1-(2-Fluorophenyl)-4-Methoxypiperidin-4-yl (Fpmp) Protecting Group in the Solid-Phase Synthesis of Oligo- and Poly-ribonucleotides," J. Chem. Soc. Perkin Trans. 1993, pp. 43-55	
Ho	9	HIMANSHU RASTOGI, et al. "A New 2'-Hydroxyl Protecting Group for the Automated Synthesis of Oligoribonucleotides," Nucleic Acids Research, 1995, Vol. 23, No. 23, pp. 4872-4877	
Ho	10	OSAMU SAKATSUME, et al. "Synthesis and Properties for Non-hammerhead RNA Using 1-(2-Chlorethoxy)- Ethyl Group for the Protection of 2'-Hydroxyl Function," Nueclosides & Nucleotides, 10 (1-3), 1991, pp. 141-153	
Ho	11	MITSUO SEKINE, et al. "Cyclic Orthoester Functions as New Protecting Groups in Nucleosides," J. American Chemical Society," 1983, 105, pp. 2044-2049	

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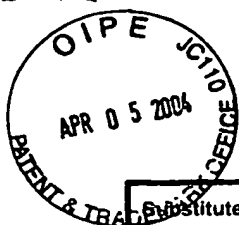
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AS	12	TOSHIKI TANAKA, et al. "Solid Phase Synthesis of Oligoribonucleotides Using 0-Nitrobenzyl Protection of 2'-hydroxyl via a Phosphite Triester Approach," Nucleic Acids Research, Vol. 14, Number 15, 1986, pp. 6265-6279			
AS	13	WU et al. (1989) "Prevention of Chain of Cleavage in the Chemical Synthesis of 2' - Silylated Oligoribonucleotides," Nucleic Acids Research, Vol. 17, No. 9. 1989, pp.3501-3517			
AS	14	STEPHEN A. SCARINGE, et al. "Chemical Synthesis of Biologically Active Oligoribonucleotides Using β -Cyanoethyl Protected Ribonucleoside Phosphoramidites," Nucleic Acids Research, Vol. 18, No. 18, pp. 5433-5441 <i>f 1990.</i>			
EXAMINER SIGNATURE		<i>Richard J. [Signature]</i>		DATE CONSIDERED	3-7-05
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